

Neovascular Age-Related Macular Degeneration Treatment Trial with Ranibizumab (MARINA)



Objective

To evaluate whether ranibizumab, an anti-VEGF antibody, improves patient outcomes and prevents vision loss in patients with neovascular age-related macular degeneration (nAMD).

Methods

Design: Multicenter RCT

Sample Size: 716

Treatment Groups:

- 238 to sham injection
- 238 to 0.3mg ranibizumab
- 240 to 0.5mg ranibizumab
- *PDT allowed if indicated*

Outcomes:

- Primary end point: Proportion of patients that lost fewer than 15 letters compared to their baseline visual acuity (VA) at 12 months
- Improvement of VA by 15 or more letters
- Mean increase in VA

Results

Point 1: Patients treated with ranibizumab exhibited less decline from baseline VA compared to sham-injections

- 94.5% of patients receiving 0.3mg and 94.6% of patients receiving 0.5mg lost fewer than 15 letters from baseline VA compared to the 62.2% that received sham injections ($P < 0.001$)

Point 2: Patients treated with ranibizumab showed improved VA.

- Roughly 1/4 of patients receiving 0.3 mg and 1/3 of patients receiving 0.5mg injections gained 15 or more letter in VA compared with 5% or less of those in the sham-injection group ($P < 0.001$)

Point 3: Mean VA with notable improvement for treatment groups

- Increase of 6.5 letters in the 0.3 mg Ranibizumab group
- Increase of 7.2 letters in the 0.5 mg Ranibizumab group
- Decrease of 10.4 letter decrease in the sham-injection group

TLDR: Treatment of nAMD with intravitreal ranibizumab prevented vision loss and improved mean VA in patients with minimally classic or occult choroidal neovascularization.